AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listing, of claims in the application:

Listing of Claims:

1. (Currently Amended) A pharmaceutical composition comprising as an active ingredient at least one <u>solid</u> olanzapine polymorph selected from Form III olanzapine, Form IV olanzapine, Form V olanzapine, and salts and mixtures thereof; and

one or more pharmaceutically acceptable carriers, excipients or diluents;

wherein Forms III, IV and V olanzapine are olanzapine polymorphs having typical x-ray powder diffraction patterns represented by the following interplanar spacings:

FORM-III	FORM-IV	FORM-V
d-spacing (Å)	d-spacing (Å)	d-spacing (Å)
10.3156	9.9487	10.5932
7.1713	8.5074	10.2170
6.5014	8.2103	9.9503
5.5165	4.8172	8.5259
4.8541	4.7114	7.1016
4.5578	4.6122	6.0731
4.4938	4.5282	5.2041
4.4536	4.2340	4.9856
4.2588	4.0901	4.8153
3.9898	3.7574	4.7514
3.7288	3.6989	4.5302
3.5626		4.4714
3.0262		4.2271
		4.1307

3.9880
3.7763
3.7167
3.5315.

- 2. (Original) The pharmaceutical composition according to claim 1, wherein the olanzapine polymorph is Form III olanzapine.
- 3. (Original) The pharmaceutical composition according to claim 2, wherein the Form III olanzapine is further characterized by substantially the following x-ray powder diffraction pattern, wherein d represents the interplanar spacing and I/I₁ represents the typical relative intensities:

d-spacing (Å)	I/I_1
10.3156	100
7.1713	16
6.5014	17
5.5165	24
4.8541	46
4.5578	24
4.4938	38
4.4536	36
4.2588	49
3.9898	52
3.7288	42
3.5626	25
3.0262	18.

4. (Original) The pharmaceutical composition according to claim 2, wherein the Form III olanzapine is further characterized by having an infrared spectrum having absorbances at the following wavenumbers:

- 5. (Original) The pharmaceutical composition according to claim 1, wherein the olanzapine polymorph is Form IV olanzapine.
- 6. (Original) The pharmaceutical composition according to claim 5, wherein the Form IV olanzapine is further characterized by substantially the following x-ray powder diffraction pattern, wherein d represents the interplanar spacing and I/I₁ represents the typical relative intensities:

I/I_1
83
15
17
100
41
35
33
29
32
23

7. (Original) The pharmaceutical composition according to claim 5, wherein the Form IV olanzapine is further characterized by having an infrared spectrum having absorbances at the following wavenumbers:

- 8. (Original) The pharmaceutical composition according to claim 1, wherein the olanzapine polymorph is Form V olanzapine.
- 9. (Original) The pharmaceutical composition according to claim 8, wherein the Form V olanzapine is further characterized by substantially the following x-ray powder diffraction pattern, wherein d represents the interplanar spacing and I/I₁ represents the typical relative intensities:

d-spacing (Å)	I/I_1
10.5932	17
10.2170	100
9.9503	57
8.5259	22
7.1016	17
6.0731	17
5.2041	19

4.9856	20
4.8153	62
4.7514	34
4.5302	24
4.4714	51
4.2271	91
4.1307	40
3.9880	31
3.7763	10
3.7167	. 62
3.5315	22.

10. (Original) The pharmaceutical composition according to claim 8, wherein the Form V olanzapine is further characterized by having an infrared spectrum having absorbances at the following wavenumbers:

11. (Currently Amended) A pharmaceutical composition containing as an active ingredient at least one <u>solid</u> olanzapine polymorph selected from Form III olanzapine, Form IV olanzapine, Form V olanzapine, and salts and mixtures thereof;

wherein Forms III, IV and V olanzapine are olanzapine polymorphs having typical x-ray powder diffraction patterns represented by the following interplanar spacings:

FORM-III	FORM-IV	FORM-V
d-spacing (Å)	d-spacing (Å)	d-spacing (Å)
10.3156	9.9487	10.5932
7.1713	8.5074	10.2170
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4.8541	4.7114	7.1016
4.5578	4.6122	6.0731
4.4938	4.5282	5.2041
4.4536	4.2340	4.9856
4.2588	4.0901	4.8153
3.9898	3.7574	4.7514
3.7288	3.6989	4.5302
3.5626		4.4714
3.0262		4.2271
		4.1307
		3.9880
		3.7763
		3.7167
		3.5315.

12. (Previously Presented) A method of treating a patient having a psychotic condition or mild anxiety comprising administering a therapeutically effective amount of at least one olanzapine polymorph to said patient;

wherein the at least one olanzapine polymorph is selected from Form III olanzapine, Form IV olanzapine, Form V olanzapine, and salts and mixtures thereof; and

wherein Forms III, IV and V olanzapine are olanzapine polymorphs having typical x-ray powder diffraction patterns represented by the following interplanar spacings:

FORM-III	FORM-IV	FORM-V
d-spacing (Å)	d-spacing (Å)	d-spacing (Å)
10.3156	9.9487	10.5932
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4.4938	4.5282	5.2041
4.4536	4.2340	4.9856
4.2588	4.0901	4.8153
3.9898	3.7574	4.7514
3.7288	3.6989	4.5302
3.5626		4.4714
3.0262		4.2271
		4.1307
		3.9880
		3.7763
		3.7167
		3.5315.

- 13. (Original) The method according to claim 12, wherein the olanzapine polymorph is Form III olanzapine.
- 14. (Original) The method according to claim 12, wherein the olanzapine polymorph is Form IV olanzapine.
- 15. (Original) The method according to claim 12, wherein the olanzapine polymorph is Form V olanzapine.
- 16. (Previously Presented) A method of treating a patient having a psychotic condition selected from schizophrenia and schizophreniform disorders, acute mania, Bipolar I Disorder, psychotic mood disorder and psychosis associated with Alzheimer's disease comprising administering a therapeutically effective amount of at least one olanzapine polymorph to said patient;

wherein the at least one olanzapine polymorph is selected from Form III olanzapine, Form IV olanzapine, Form V olanzapine, and salts and mixtures thereof; and

wherein Forms III, IV and V olanzapine are olanzapine polymorphs having typical x-ray powder diffraction patterns represented by the following interplanar spacings:

FORM-III	FORM-IV	FORM-V
d-spacing (Å)	d-spacing (Å)	d-spacing (Å)
10.3156	9.9487	10.5932
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5.5165	4.8172	8.5259
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4.5578	4.6122	6.0731
4.4938	4.5282	5.2041
4.4536	4.2340	4.9856

4.2588	4.0901	4.8153
3.9898	3.7574	4.7514
3.7288	3.6989	4.5302
3.5626		4.4714
3.0262		4.2271
		4.1307
		3.9880
		3.7763
		3.7167
		3.5315.

- 17. (Original) The method according to claim 16, wherein the olanzapine polymorph is Form III olanzapine.
- 18. (Original) The method according to claim 16, wherein the olanzapine polymorph is Form IV olanzapine.
- 19. (Original) The method according to claim 16, wherein the olanzapine polymorph is Form V olanzapine.
- 20. (Previously Presented) The composition of claim 1, in the form of a suspension, emulsion, solid capsule, granular formulation or powdered formulation.